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Sheet 1 of 1

Complete if Known

Application Number 09/529,053-Conf. #1413
Filing Date April 6, 2000
First Named Inventor James W. Waldman
Art Unit 1617
Examiner Name S. Wang
Attorney Docket Number 28385/35415

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Document Number Number-Kind Code ² (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear

FOREIGN PATENT DOCUMENTS

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NON PATENT LITERATURE DOCUMENTS

Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
	E1	Ashour et al., 5-(m-Benzyloxybenzyl) barbituric Acid Acyclonucleoside, a Uridine Phosphorylase Inhibitor, and 2',3',5'-Tri-O-Acetyluridine, a Prodrug of Uridine, as Modulators of Plasma Uridine Concentration. Biochemical Pharmacology, Vol. 15, pp. 1601-1611 (1996)	
	E2	Hidalgo et al., Phase I and Pharmacologic Study of PN401 and Fluorouracil in Patients with Advanced Solid Malignancies, Journal of Clinical Oncology, Vol. 18, No. 1 pp. 167-177 (2000)	
	E3	Kelsen et al., Phase I Trial and PN401, an Oral Prodrug of Uridine, to Prevent Toxicity from Fluorouracil in Patients with Advanced Cancer, Journal of Clinical Oncology, Vol. 15, No. 4 pp. 1511-1517 (1997)	
	E4	Van Groeningen et al., Reversal of 5-fluorouracil-induced toxicity by oral administration of uridine, Annals of Oncology 4:317-320 (1993)	
	E5	Van Groeningen et al., Clinical and Pharmacologic Study of Orally Administered Uridine, Journal of the National Cancer Institute, Vol. 83, No. 6, pp. 437-441 (1991)	
	E6	The Merck Index Eleventh Edition, Merck & Co., NJ (1989), pp. 437, 1480, 1554.	
	E7	Hawley's Condensed Chemical Dictionary Thirteenth Edition, Van Nostrand Reinhold, New York, NY (1997), pp. 329, 1105, 1159.	

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